

K770938 TINNITUS AID, MODEL S244Jun 28, 1977
36 days to decisionK770938 · Product code: **KLW** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k770938/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Masker, Tinnitus (KLW)
Date received	May 23, 1977
Decision date	Jun 28, 1977
Days to decision	36 days
Third-party review	No

APPLICANT

Company	Vicon Instrument Co.
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1976-1981

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Device record: <https://www.510kdatabase.net/k770938/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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